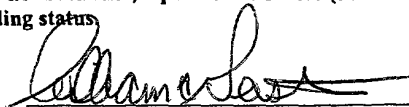


414 Rec'd PCT/PTO 22 JUN 2000

FORM PTO-1390 (REV 12-29-99)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		ATTORNEY'S DOCKET NUMBER	
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371				STUDIEN 272-KGB	
				U.S. APPLICATION NO. (If known, see 37 CFR 1.5) 09/582116	
INTERNATIONAL APPLICATION NO. PCT/EP98/08214		INTERNATIONAL FILING DATE 15 Dec. 1998 (15.12.98)		PRIORITY DATE CLAIMED 23 Dec. 1997 (23.12.97)	
TITLE OF INVENTION METHOD FOR THE COMBINATORIAL DEVELOPMENT OF MATERIALS BY USING HEAT DIFFERENCE IMAGES					
APPLICANT(S) FOR DO/EO/US Wilhelm MAIER; Arnold HOLZWARTH					
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:					
<ol style="list-style-type: none"> 1. <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371. 2. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371. 3. <input checked="" type="checkbox"/> This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1). 4. <input checked="" type="checkbox"/> A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date. 5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371(c)(2)) <ol style="list-style-type: none"> a. <input type="checkbox"/> is transmitted herewith (required only if not transmitted by the International Bureau). b. <input checked="" type="checkbox"/> has been transmitted by the International Bureau. c. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US). 6. <input checked="" type="checkbox"/> A translation of the International Application into English (35 U.S.C. 371(c)(2)). 7. <input type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3)) <ol style="list-style-type: none"> a. <input type="checkbox"/> are transmitted herewith (required only if not transmitted by the International Bureau). b. <input type="checkbox"/> have been transmitted by the International Bureau. c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired. d. <input type="checkbox"/> have not been made and will not be made. 8. <input type="checkbox"/> A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)). 9. <input type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)). 10. <input type="checkbox"/> A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)). 					
Items 11. to 16. below concern document(s) or information included:					
<ol style="list-style-type: none"> 11. <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98. 12. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included. 13. <input checked="" type="checkbox"/> A FIRST preliminary amendment. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment. 14. <input type="checkbox"/> A substitute specification. 15. <input type="checkbox"/> A change of power of attorney and/or address letter. 16. <input checked="" type="checkbox"/> Other items or information: <ol style="list-style-type: none"> a) copy of cover page from published application WO 99/34206 b) five (5) sheets of drawings (Figs. 1-7), attached to English translation of application 					

09/582116 U.S. APPLICATION NO. (37 CFR 1.53) INTERNATIONAL APPLICATION NO. PCT/EP98/08214		ATTORNEY'S DOCKET NUMBER STUDIEN 272-KGB	
17. <input checked="" type="checkbox"/> The following fees are submitted: BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)): Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO \$970.00 International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO \$840.00 International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO \$690.00 International preliminary examination fee paid to USPTO (37 CFR 1.482) but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$670.00 International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(1)-(4) \$96.00 ENTER APPROPRIATE BASIC FEE AMOUNT =		CALCULATIONS PTO USE ONLY 	
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).		\$ 840.00	
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE
Total claims	15 - 20 =	0	X \$18.00
Independent claims	1 - 3 =	0	X \$78.00
MULTIPLE DEPENDENT CLAIM(S) (if applicable)		+ \$260.00	\$ 260.00
TOTAL OF ABOVE CALCULATIONS =		\$ 1,100.00	
Reduction of 1/2 for filing by small entity, if applicable. A Small Entity Statement must also be filed (Note 37 CFR 1.9, 1.27, 1.28).		\$ 550.00	
SUBTOTAL =		\$ 550.00	
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).		\$	
TOTAL NATIONAL FEE =		\$ 550.00	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property +		\$	
TOTAL FEES ENCLOSED =		\$	
		Amount to be:	\$
		refunded	\$
		charged	\$
a. <input type="checkbox"/> A check in the amount of \$_____ to cover the above fees is enclosed. b. <input checked="" type="checkbox"/> Please charge my Deposit Account No. <u>14-1263</u> in the amount of \$ <u>14-1263</u> to cover the above fees. A duplicate copy of this sheet is enclosed. c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. <u>14-1263</u> . A duplicate copy of this sheet is enclosed.			
NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.			
SEND ALL CORRESPONDENCE TO: Kurg G. Briscoe NORRIS, McLAUGHLIN & MARCUS, P.A. 660 White Plains Road Tarrytown, New York 10591		<div style="text-align: center;">  SIGNATURE: <u>William C. Gerstenzang</u> NAME <u>27,552</u> REGISTRATION NUMBER </div>	

STUDIEN 272-KGB
PCT/EP98/08214

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant(s) : Wilhelm MAIER; Arnold HOLZWARTH
Serial No. : To Be Assigned
Filed : Herewith
For : METHOD FOR THE COMBINATORIAL
DEVELOPMENT OF MATERIALS BY USING HEAT
DIFFERENCE IMAGES
Art Unit : To Be Assigned
Examiner : To Be Assigned

June 21, 2000

BOX PCT
Hon. Assistant Commissioner For Patents
Washington, D.C. 20231

PRELIMINARY AMENDMENT

Sir:

In advance of prosecution, the Examiner is respectfully requested to
amend the application as follows and consider the following remarks:

IN THE CLAIMS

Claim 1 (amended). A method for the comparative determination of
the heat changes caused by physical or chemical processes, [characterized in
that] wherein a difference image is recorded using an infrared camera, which
image corresponds to a subtraction of the infrared emission recorded prior to the

beginning of the processes from the infrared emission recorded during the course of the processes.

Claim 6 (amended). The method according to [any of claims 2 to 4] Claim 2, 3 or 4, wherein said library plate consists of a material having low infrared reflectivity.

Claim 8 (amended). The method according to [any of claims 2 to 4] Claim 2, 3 or 4, wherein said library plate is coated with a film having low infrared reflectivity.

Claim 9 (amended). The method according to [any of claims 2 to 7] Claim 2, 3, 4 or 5, wherein the region of the surface of the library plate not occupied by catalysts or materials is coated with a non-wetting film.

Claim 13 (amended). The method according to [any of claims 2 and 4-12] Claim 2, 4, 5, 10 or 11, wherein the catalysts are within a reactor under reaction conditions and are recorded by the externally provided infrared camera through an infrared-transparent window.

Claim 14 (amended). The method according to [any of claims 1 to 13] Claim 1, 2, 3, 4, 5, 10 or 11, wherein the infrared emission is recorded through a

wavelength-specific infrared filter.

REMARKS

This Preliminary Amendment is being filed to conform to conventional format, and to eliminate the dependency of multiple dependent claims on other multiple dependent claims.

Favorable action is respectfully solicited.

ADDITIONAL FEE

Please charge any insufficiency of fees, or credit any excess, to Deposit Account No. 14-1263.

P.A.

Respectfully submitted,
NORRIS, McLAUGHLIN & MARCUS,

By 

William C. Gerstenzang
Reg. No. 27,552
for Kurt G. Briscoe
Reg. No. 33,171

660 White Plains Road
Tarrytown, NY 10591-5144
(914) 332-1700

I hereby certify that this paper is being deposited with the United States Postal Service as Express Mail, Label No. EH976287793US, addressed to: Hon. Asst. Commissioner of Trademarks, 2900 Crystal Drive, Arlington, Virginia 22202-3513.

Norris McLaughlin & Marcus, P.A.

By: 

Date: 

S/PRB

09/582116

430 Rec'd PCT/PTO 22 JUN 2000

SMB

Method for the Combinatorial Development of Materials
by Using Heat Difference Images

The present invention relates to a method for the comparative determination of properties of the materials of a combinatorial library using an infrared camera which records the heat changes of chemical or physical processes in the form of a difference image. The essential features of the invention have already been described in the application DE 197 57 754.7 of December 23, 1997. The present application contains an extension of the method by the use of selective IR filters, its use for the general characterization of material properties, its use for homogeneous catalysts, and an improved embodiment with library plates of low IR reflectivity, such as slabs of slate.

Prior Art

The narrowest bottleneck in the development of new materials is the discovery of suitable new leading structures. In particular, this applies to new catalysts. Despite the wide-spread use of heterogeneous and homogeneous catalysts in chemical technology and the related extensive research efforts, the detailed mechanism of action of heterogeneous and homogeneous catalysts under industrial reaction conditions is not sufficiently known. Therefore, one has to rely on empirical methods for selecting appropriate catalyst materials for each application and finding reaction conditions under which a desired catalytic activity and selectivity occurs. Classical test methods for catalysts require that each candidate catalyst material is individually tested under the reaction conditions, which is a very expensive and also time-consuming procedure in view of the immeasurable abundance of possible materials. A promising solution is offered in the field of combinatorial materials sciences for making a reasonable

- 2 -

preselection of heterogeneous and homogeneous catalyst materials for technically interesting reactions and limiting the test method to a few promising materials. While heterogeneous catalysts are predominantly important to the production of mass products and fine chemicals, homogeneous catalysts are dominating in the production of optically pure active substances in the pharmaceutical field and in tacticity-controlled polymerization.

In the so-called combinatorial methods, a large number of samples is applied to a support within a very small space. Then, using readily available physical measuring quantities associated with the desired property of the material, a fast and reliable test is performed for this desired property on every sample.

In this way, new magnetoresistant materials [B. Gabriel, H. Chang, X. Sun, P.G. Schultz, X.-D. Xiang, "A Class of Cobalt Oxide Magnetoresistance Materials Discovered with Combinatorial Synthesis", *Science*, 270, 273-275, 1995], new superconducting materials [X.-D. Xiang, X. Sun, G. Briceno, Y. Lou, K.-A. Wang, H. Chang, W.G. Wallace-Freedman, S.-W. Chen, P.G. Schultz, "A Combinatorial Approach to Materials Discovery", *Science*, 268, 1738-1740, 1995] and new luminescent materials [X.-D. Sun, K.-A. Wang, Y. Yoo, W.G. Wallace-Freedman, C. Gao, X.-D. Xiang, P.G. Schultz, "Solution-Phase Synthesis of Luminescent Materials Libraries", *Adv. Mater.* 9, 1046-1049, 1997] have already been discovered.

Even in the field of combinatorial chemistry of heterogeneous catalysts, corresponding publications already exist [F.C. Moates, M. Somani, J. Annamalai, J.T. Richardson, D. Luss, R.C. Willson, "Infrared Thermographic Screening of Combinatorial Libraries of Heterogeneous Catalysts", *Ind. Eng. Chem. Res.* 35, 4801-4803, 1996; R.C. Willson, "Catalyst Testing Process and Apparatus", U.S. Patent: PCT/US 97/02756, International Patent: WO 97/32208]. The authors describe an experiment in which catalytic activity is detected from the accompanying **heat of reaction** using an infrared camera. However, the detection sensitivity of the configuration described by the authors was relatively low so that only the extremely exothermic hydrogen-oxygen reaction (14 kJ/g) was detected. This is not a reaction of technical importance. To effectively

- 3 -

examine catalysts for technically important reactions, above all, a higher detection sensitivity of the measuring configuration is necessary, and evidence must be provided, that heats of reaction can be registered at low temperature with different reactions and with minute amounts of catalyst. The recognition of the catalytic activity of polymer-bound catalysts in solution was demonstrated on coded beads (0.5 mm) [S.J. Taylor, J.P. Morken, Science 280 (1998) 268]. The density of the solution must be higher than the density of the beads because the latter can only be observed when afloat on the surface. Drawbacks are the expensive coding technique and the necessity of using a solvent of higher density than that of the floating fixed-bed catalysts. In addition, there is a problem in using uncorrected infrared images, i.e., the fact that due to different emission properties, highly sensitive infrared cameras indicate the presence of different materials on a surface when no reactions take place so that the practical utility of such methods is highly limited.

The reliable preparation of catalyst libraries is another important aspect. Well known are combinatorial libraries, although largely restricted to the field of pharmaceutical applications, the generation of which can partly be transferred to the preparation of libraries for developing new materials as well (B. Posner et al., Trends in Biochemical Science 19 (1994) 145). One method which is probably the most reliable and most widely applicable one for the preparation of combinatorial libraries of materials is the combination of ink-jet printing technology with computer control which allows to produce the new materials dot by dot by mixing solutions in the nanoliter range and to convert them to the solids to be examined by a well-defined posttreatment [X.-D. Sun, K.-A. Wang, Y. Yoo, W.G. Wallace-Freedman, C. Gao, X.-D. Xiang, P.G. Schultz, "Solution-Phase Synthesis of Luminescent Materials Libraries", Adv. Mater. 9, 1046-1049, 1997]. However, the problem here is the use of suitable synthetic conditions which allow the production of defined catalyst and other materials from the mixing of nanoliter solutions.

- 4 -

Description of the invention

We have now found that minute heat changes occurring in the course of chemical or physical processes can be reliably visualized with an infrared camera with spatial resolution when a difference heat image is recorded which results from the subtraction of the infrared emission intensities recorded prior to the beginning of the reaction from the intensities obtained under reaction conditions. Suitable chemical or physical processes with heat changes include chemical reactions, phase transitions, conversions, sorption processes (physisorption, chemisorption, adsorption, desorption), absorption processes (absorption of molecules or electromagnetic radiation), magnetic induction. Thus, the heat changes of the physisorption of molecules from the gas-phase can be utilized for recognizing porosities in materials. The heat of catalyzed reactions serves for recognizing the catalytic activity of materials. The heat changes of phase transitions serve for recognizing changes in substances. The heat changes associated with surface reactions, such as the formation of carbides or nitrides by reactions of alloys or mixed oxides with methane or ammonia serve for recognizing conversions. The heat changes associated with chemisorption serves for recognizing surface reactivity, such as acidic or basic sites. The heat changes associated with the absorption of electromagnetic radiation characterize particular absorption properties, such as ultraviolet absorption, X-ray absorption, or absorption of visible light. Magnetic properties are recognized by the heating brought about by applying a magnetic field. All these properties are of particular interest when specific materials are significantly different from the bulk of other materials and thus represent potential novel lead structures.

In particular, the method of interest for the development of new catalysts by the use of combinatorial libraries. Surprisingly, a suitable method for the preparation of combinatorial catalyst libraries with heterogeneous catalysts is the sol-gel method, as partially described in DE-A-195.45042.6. In addition, we have found that the heat of homogeneously catalyzed reactions can also be reliably visualized with this technique. The use of heat difference images also allows for a temperature calibration and thus a reliable

- 5 -

assignment of the actual temperature increase at the active catalyst. This is of great importance to the evaluation of relative activities and quantifications of the catalyst performances.

The catalytic reactions are preferably conducted under exactly controlled conditions in a hermetically sealed reactor provided with an infrared-transparent window to enable recording with a camera.

Catalyst libraries consisting of catalyst components in the form of metal oxides and/or mixed metal oxides are prepared by first distributing the precursors of the catalyst components to be examined as an aqueous or alcoholic solution of silicon or metal compounds in the form of their alkoxy derivatives, mixed alkoxy derivatives, alkoxyoxo or acetylacetonate derivatives or in the form of their halides, nitrates, citrates or other carboxylates over the surface of a plate consisting of, for example, slate, metal or steel, glass, ceramic or plastic, e.g., in recesses formed by drilling into the plate. In particular, the catalyst component can be carbides, nitrides and zeolites. The library plates are subsequently dried and calcined. When the catalyst library is prepared, it may happen, due to the surface properties of the library plate, that the sol creeps out of the bores onto the surface of the library plate. This can be prevented by coating the library plate with a non-wetting infrared-transparent film in the regions not occupied by catalysts, for example, if the coating is applied before making the bores.

Not only solids, but also molecules in gases emit infrared radiation. The intensity of this thermal emission increases with temperature. Especially by reflections of this emitted radiation at reactor walls and at the surface of the catalyst library, the false impression of inhomogeneous increases of temperature can be evoked. Reflections at the catalyst library are particularly inconvenient because they are directly recorded by the infrared camera. For this reason, the radiation properties of the support material of the catalyst library are of critical importance. Therefore, a material having a particularly low reflectivity, close to that of a black body is particularly suitable as a support material for the catalyst library. This effect may also be achieved by a suitable coating. The interior walls of the reactor should also reflect as little infrared radiation as possible. This can

- 6 -

be achieved, for example, by appropriate antireflection coatings. In addition, the support material for the catalyst library should have as low as possible a thermal conductivity in order that heat developed at the catalyst sites during the catalytic reaction is not dissipated too fast and an increase in temperature as high as possible can be achieved.

Wavelength ranges in which the gaseous reactants and products display a particularly high infrared activity are unfavorable for IR-thermographic temperature measurements since interfering effects from the absorption and emission by the gases are more prevailing in this case. Such wavelength ranges can be reduced using a wavelength-specific infrared filter. For gaseous hydrocarbons, we have found it convenient to reduce the particularly intensive CH vibrations with such a filter.

IR thermography as described above is also suitable for recognizing the catalytic activity of enantioselective homogeneous catalysts and enzymes. By using optically pure reactants or racemates, both the relative catalytic activities of different catalysts and their enantioselectivity can be recognized by IR thermography. Reactions examined include the enantioselective acylation of 1-phenylethanol with lipase [M.T. Reetz, A. Zonta, J. Simpelkamp, *Angew. Chem.* 107 (1995) 373-376] and the hydrolysis of chiral epoxides with Mn, Co and Cr salen complexes [M. Tokunaga, J.F. Larrow, F. Kakiuchi, E.N. Jacobsen, *Science* 277 (1997) 936-9389] in a small combinatorial library of 9 reaction solutions. Using the IR-thermographic difference images, both the relative activities of the catalysts and their enantioselectivity could be simultaneously visualized.

The method can of course be applied to many other types of reactions, such as selective hydrogenations, selective oxidations, esterifications, pericyclic reactions, halogenations, dehalogenations, hydrogenolyses, hydrations, dehydrations, condensations, enantioselective reactions, polymerization reactions, polycondensation reactions, oxidative couplings, and selective oxidations of olefins and alkanes, especially with air, H₂O₂ or ozone with the selective formation of epoxides, ketones, aldehydes, alcohols, carboxylic acids or anhydrides. Heats of such reactions in liquid phase can also be

- 7 -

visualized very simply and effectively using an infrared camera (Maier, Holzwarth, DE 19757 754.7).

Examples

Construction of the camera, correction of the detector and temperature calibration:

In the following Examples, an AIM AEGALS infrared camera was employed with a PtSi (platinum/silicon) detector with 256 times 256 pixels as described in the following. The camera was connected with a computer. The software (version 1.40 of July 4, 1997) allowed to subtract the background in such a way that only an increase in temperature relative to the starting temperature is indicated in each pixel.

Directly before the entrance window of the detector, a 3.6 μm infrared cut-off filter was applied which is opaque to infrared radiation having wavelengths of shorter than 3.6 μm . The inhomogeneity of the detector was corrected for by a two-point correction. Thus, IR images of the library were recorded 5 °C below and above the reaction temperature (hydrogenation: 100 °C; oxidation: 350 °C; lipase catalysis: 30 °C; metal salene catalysis: 27 °C). After reaching the reaction temperature prior to the start of the reaction, another infrared image of the catalyst library was recorded. This image was then subtracted from all following infrared images (offset) so that only temperature changes due to catalytic activity at the catalyst sites became visible through corresponding color changes. The temperature calibration was effected by recording in the catalyst library in the reactor at 6 temperatures, followed by fitting the individual pixels using a quadratic polynomial, a function provided for in the camera software.

- 8 -

Example 1:Combinatorial test method for screening the catalytic activity of materials from a libraryConstruction of the reactor:

When the reactor was constructed, particular care was taken that a temperature distribution as homogeneous as possible within the reactor was achieved. The reactor essentially consists of two portions, the reactor head with an infrared-transparent barium fluoride window for observing the catalyst library, and the reactor block, which consists of solid steel. The reactor head with the infrared-transparent barium fluoride window forms an angle of 10° . The heating elements are in the lower portion of the reactor. They are arranged in such a way that the complete reactor block and reactor head are heated up as homogeneously as possible. The gas inlets were led from below through the reactor block into the reaction chamber with the catalyst library in order to preheat the gaseous educts to the same temperature as prevails at the catalyst library prior to the reaction at the catalyst sites. The gas mixture was fed through a multitude of small holes arranged around the circular catalyst library in order to achieve as homogeneous as possible a distribution of the gases in the chamber.

Preparation of the silica sols:

Silica and titania sols were prepared and pipetted into appropriate bores on a slate library plate using a microliter pipette. After the solvent was evaporated, the whole library was calcined. The plate was inserted into a special reactor, designed for observation with the infrared camera (see above). Then, the catalysts on the plate were activated by tempering at 300°C in a hydrogen flow for three hours. Then, experiments for the hydrogenation of 1-hexyne were performed at 100°C . Finally, the experiments on the oxidation of isooctane and toluene with synthetic air were performed at 350°C .

- 9 -

Silica sols were prepared by the standard protocol of Klein et al. [S. Klein, S. Thorimbert, W.F. Maier, J. Catal. 163 (1996) 476-488]. Tetraethoxy orthosilicate served as a silica precursor. The following compounds were dissolved in the sol as precursors of the other elements: PdCl_2 , Na_2PtCl_6 , $\text{IrCl}_4 \cdot \text{H}_2\text{O}$, $\text{RuCl}_3 \cdot \text{H}_2\text{O}$, $\text{CoCl}_2 \cdot 6\text{H}_2\text{O}$, $\text{Fe}(\text{acac})_3$, $\text{Mn}(\text{acac})_3$, $(\text{iPrO})_3\text{VO}$, $\text{CrCl}_3 \cdot 6\text{H}_2\text{O}$, $\text{Cu}(\text{acac})_2$, $\text{Ni}(\text{acac})_2$, ZnCl_2 , $\text{Pd}(\text{acac})_2$, $\text{RhCl}_3 \cdot 3\text{H}_2\text{O}$.

Preparation of the titania sols:

With constant stirring, 1 ml of titanium isopropoxide (3.36 mmol) was dissolved in 3.33 ml of dry ethanol. After 30 minutes of stirring, 8.33 ml of 8 N HCl was added. After another 5 minutes, 46.7 ml of 12 N HCl was slowly added in the course of 20 min. Then, 833 ml of ethanol was added. Finally, 833 ml of an ethanolic solution of the metal compound was added.

Preparation of the catalyst library:

As the material for the catalyst library, slate was selected due to its low thermal conductivity and its low reflectivity for infrared radiation. The circular library plate (diameter 5 cm) had 69 holes with diameters of 1.5 mm and a depth of 0.6 mm.

The catalyst library was prepared by pipetting microliter amounts of the silica and titania sols into the designated bores on the slate plate. In order to obtain comparable amounts of catalyst material, 1.5 μl was pipetted in the case of the silica sols (corresponding to 192 μg of silica), and 5 μl was pipetted in the case of the titania sols (corresponding to 182 μg of titania). After evaporating the solvent, the plate was calcined as in Example 1 (temperature program: (1) from room temperature to 65 °C at 1 °C/min, (2) maintain temperature at 65 °C for 30 min, (3) from 65 °C to 250 °C at 1 °C/min, (4) maintain temperature at 250 °C for 3 h, (5) cool down to room temperature). Figure 1 shows the design and chemical compositions of the library.

- 10 -

Hydrogenation of 1-hexyne:

The hydrogenation of 1-hexyne was performed by passing hydrogen at a temperature of 100 °C through the reactor at a flow rate of 20 ml/min. 1-Hexyne was evaporated into the gas flow. The concentration of 1-hexyne in the gas phase was 0.2478 g/l.

Figure 2 shows an infrared image of the catalyst library during the reaction, the allocation of the library can be seen in Figure 1. A particularly high activity is exhibited by catalyst materials Pd₅Si, Pd₁Si, Pt₅Si and Pt₁Si.

Oxidation of isooctane with synthetic air:

The oxidation of isooctane was performed by passing synthetic air at a temperature of 350 °C through the reactor at a flow rate of 20 ml/min. Isooctane was evaporated into the gas flow. The concentration of isooctane in the gas phase was 0.2381 g/l. Figure 3 shows an infrared image of the catalyst library during the reaction. A particularly high activity is exhibited by catalyst materials Pt₂Ti, Pt₁Ti, V₅Ti, Pd₁Ti, Ru₅Ti, Cu₅Ti.

Oxidation of toluene with synthetic air:

The oxidation of toluene was performed by passing synthetic air at a temperature of 350 °C through the reactor at a flow rate of 20 ml/min. Toluene was evaporated into the gas flow. The concentration of toluene in the gas phase was 0.1489 g/l. Figure 4 shows an infrared image of the catalyst library during the reaction. A particularly high activity is exhibited by catalyst materials Pt₂Ti, Pt₁Ti, V₅Ti, Cu₅Ti, Pt₅Ti, Pt₅Si.

Example 2:

Combinatorial test method for the simultaneous screening for activity and enantioselectivity of homogeneous catalysts

Catalytic reactions in the liquid phase were performed in the wells of a modified microtitration plate. This plate was attached onto a thermally controlled Eppendorf

- 11 -

shaker. By quickly shaking the microtitration plate, it was possible to achieve homogeneous mixing of the reaction solutions within the individual wells of the plate and to control the temperature of the solutions to particular predetermined temperatures. The adjustment of the infrared camera was done as described in the previous Example. Constant shaking was applied during the reaction. For measuring with the infrared camera, the shaker was briefly stopped. In this experiment, a library of 9 reactions were observed simultaneously.

Lipase-catalyzed enantioselective acylation of 1-phenylethanol with vinyl acetate

100 μ l each of a solution of the phenylethanol as an S enantiomer, R enantiomer or racemate in toluene was pipetted into the wells of the microtitration plate. To these solutions was added 100 μ l of a solution of vinyl acetate in toluene in an equimolar ratio. Concentrations of 0.5 M, 1 M or 2 M were respectively used. Then, infrared images of the plate with the solutions for temperature calibration, two-point correction of the detector and offset were recorded. Subsequently, the reaction was started by adding 5 mg of immobilized lipase (*Candida antarctica*, Novo SP 435) at 30 °C, and infrared images of the plate with the solutions were recorded at different times during the reaction. Figure 5 shows that the heat evolution is lower for the racemate (left column) than for the R enantiomer of 1-phenylethanol (right column) whereas the S enantiomer (middle column) is not converted by the enzyme.

Enantioselective hydrolysis of epichlorohydrin with manganese-, chromium- and cobalt-salene catalysts

Three times 300 μ l each of a solution respectively containing 600 μ mol of the catalysts (S,S)-M-salene (a: M = Mn; b: M = Cr; c: M = Co) in toluene was pipetted into the wells of the microtitration plate. Then, 78.4 μ l (1 mmol) of the epichlorohydrin was added as a racemate, S or R enantiomer. Then, infrared images of the plate with the solutions were recorded for temperature calibration, two-point correction of the detector and offset. Subsequently, the reaction was started by adding 9.9 μ l (0.55 mmol) of water at 27 °C, and images of the plate with the solutions were

- 12 -

recorded at different times during the reaction. Figure 6 shows that the manganese catalyst (a) is not active, the chromium catalyst (b) shows a detectable reaction only with the S-epichlorohydrin, while the cobalt catalyst (c) is the most active, the S enantiomer reacting better than the racemate. None of the catalysts shows a detectable reaction with the R enantiomer.

Hydrolysis of different substrates with a cobalt-salene catalyst:

Into 9 wells of the microtitration plate was pipetted 100 μ l each of a solution containing 2 μ mol of the catalyst (S,S)-Co-salene in toluene. Then, solutions of epoxides d-f were added as a racemate, S or R enantiomer (d-f = epoxides of formulae $RCHCH_2O$: d: $R = CH_2OCH_2Ph$; e: $R = Ph$; f: $R = CH_2Cl$). The solutions contained the epoxides in a concentration of 3.85 mol/l. Prior to starting the reaction by the addition of water at 27 °C, infrared images of the plate with the solutions were again recorded for temperature calibration, two-point correction of the detector and offset. After the addition of water, infrared images of the plate with the solutions were recorded at different times during the reaction. Figure 7 shows that the Co catalyst selectively reacts with the S enantiomers of epoxides a and b.

- 13 -

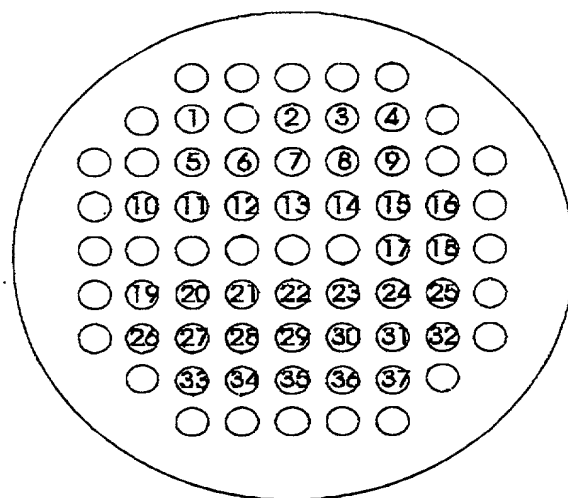
CLAIMS:

1. A method for the comparative determination of the heat changes caused by physical or chemical processes, characterized in that a difference image is recorded using an infrared camera, which image corresponds to a subtraction of the infrared emission recorded prior to the beginning of the processes from the infrared emission recorded during the course of the processes.
2. The method according to claim 1, wherein chemical reactions performed with the aid of a catalyst are observed as said chemical processes, wherein the catalysts are arranged in the form of a catalyst library over the surface of a library plate.
3. The method according to claim 1, wherein sorption processes, phase transitions or conversions at or of materials arranged on the surface of a library plate are observed as said chemical or physical processes.
4. The method according to claim 2, wherein said catalyst libraries consist of catalyst components in the form of metal oxides and/or mixed metal oxides, the precursors of which are arranged as aqueous or alcoholic solutions of silicon or metal compounds in the form of their alkoxy derivatives, mixed alkoxy derivatives, alkoxyoxo or acetylacetonate derivatives or in the form of their halides or carboxylates over the surface of the library plate, followed by drying and calcining.
5. The method according to claim 4, wherein said catalyst components are carbides, nitrides or zeolites.
6. The method according to any of claims 2 to 4, wherein said library plate consists of a material having low infrared reflectivity.

- 14 -

7. The method according to claim 6, wherein said library plate consists of slate.
8. The method according to any of claims 2 to 4, wherein said library plate is coated with a film having low infrared reflectivity.
9. The method according to any of claims 2 to 7, wherein the region of the surface of the library plate not occupied by catalysts or materials is coated with a non-wetting film.
10. The method according to claim 2, wherein said library plate contains reaction cavities comprising liquid reaction solutions with homogeneous catalysts.
11. The method according to claim 10, wherein enzymes or soluble organometallic compounds are employed as said catalysts.
12. The method according to claim 2 or 10, wherein the selectivity or the enantioselectivity of catalyzed reactions is determined on libraries.
13. The method according to any of claims 2 and 4-12, wherein the catalysts are within a reactor under reaction conditions and are recorded by the externally provided infrared camera through an infrared-transparent window.
14. The method according to any of claims 1 to 13, wherein the infrared emission is recorded through a wavelength-specific infrared filter.
15. The method according to claim 13, wherein surfaces of the reactor interior space are coated with a paint having low infrared reflectivity.

09/582116



1	Ir_1Ti	14	Ni_5Ti	27	Zn_5Si
2	Pt_1Ti	15	Rh_5Ti	28	V_5Si
3	Pt_2Ti	16	Ru_5Ti	29	Mn_3Si
4	Pt_5Ti	17	Cu_5Ti	30	Mn_5Si
5	Zn_5Ti	18	Cu_5Si	31	Fe_5Si
6	V_5Ti	19	Pd_1Si	32	Fe_{10}Si
7	Mn_3Ti	20	Pd_5Si	33	Ir_2Si
8	Mn_5Ti	21	Cr_5Si	34	Ir_5Si
9	Fe_5Ti	22	Co_5Si	35	Pt_1Si
10	Pd_1Ti	23	Ni_5Si	36	Pt_2Si
11	Pd_5Ti	24	Rh_5Si	37	Pt_5Si
12	Cr_5Ti	25	Ru_5Si		
13	Co_5Ti	26	Ti_5Si		

Figure 1/7: Design of the catalyst library of example 1: f.e. Co_5Si stands for 5 mol% cobalt oxide in 95 mol % SiO_2 , Co_5Ti stands for 5 mol% cobalt oxide in 95 mol% TiO_2

09/582116

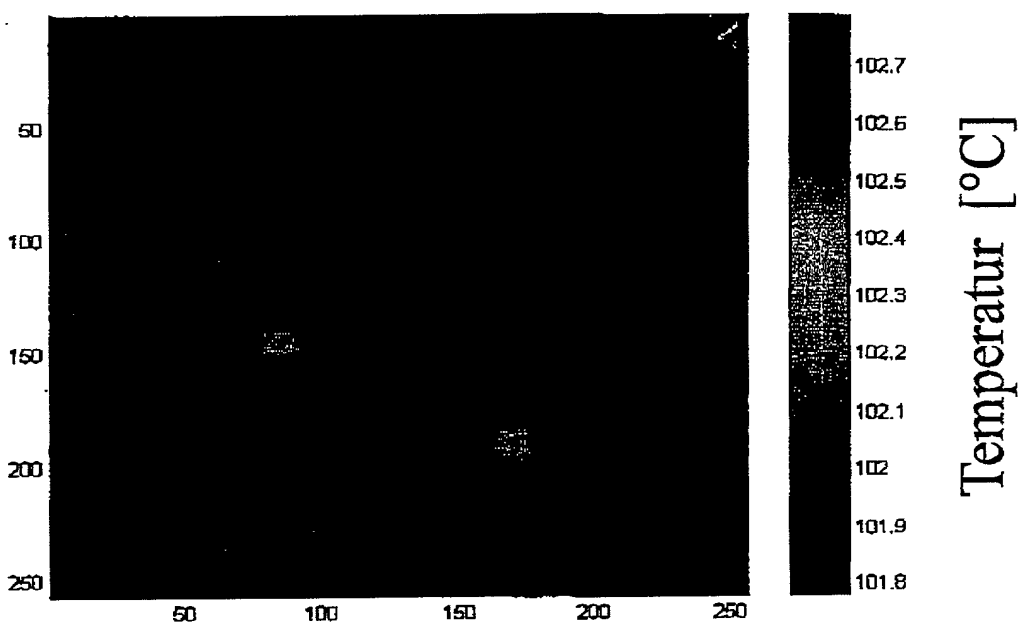


Figure 2/7: IR-image with temperature scale (right) during hydrogenation of 1-hexene at 100 °C

09/582116

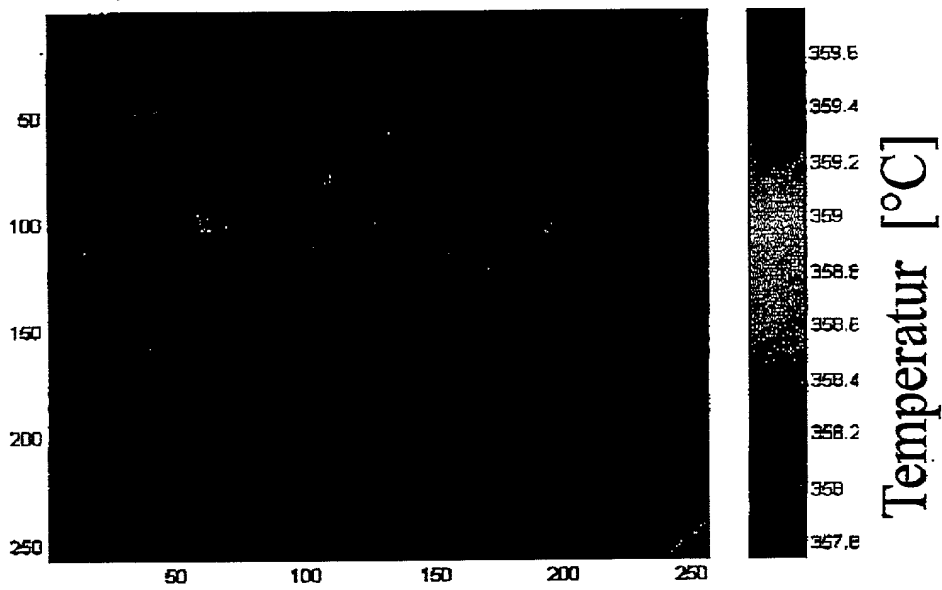


Figure 3/7: IR-image with temperature scale (right) during oxidation of isooctane with synthetic air 350 °C

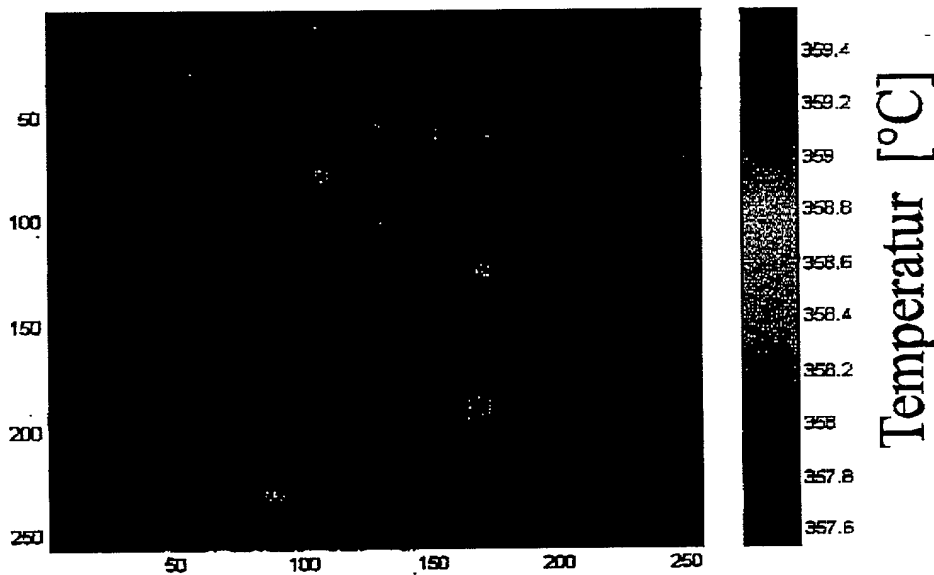


Figure 4/7: IR-image with temperature scale (right) during oxidation of toluene with synthetic air 350 °C

09/582116

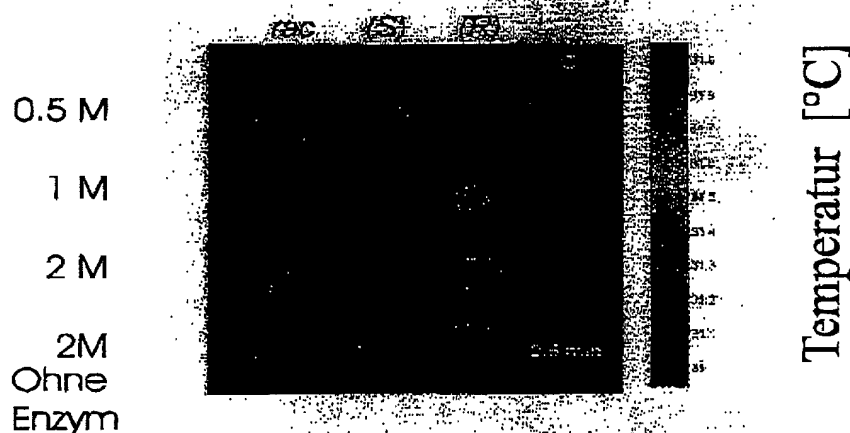


Figure 5/7: IR-image with temperature scale (right) during the lipase catalysed enantioselective acylation of 1-phenylethanol with vinylacetate at 30 °C

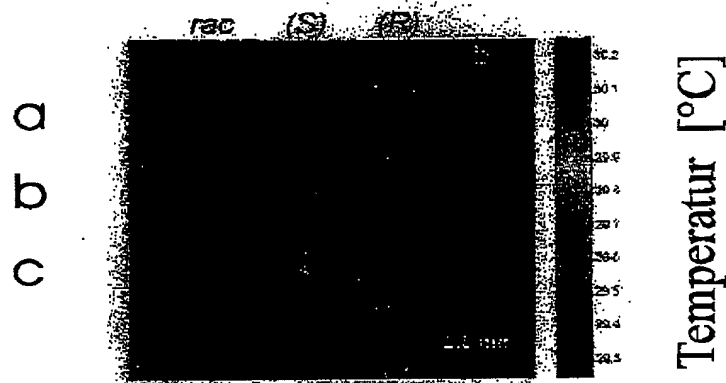


Figure 6/7: IR-image with temperature scale (right) during the enantioselective hydrolysis of epichlorohydrine with various metal-Salen catalysts at 27 °C

09/532116

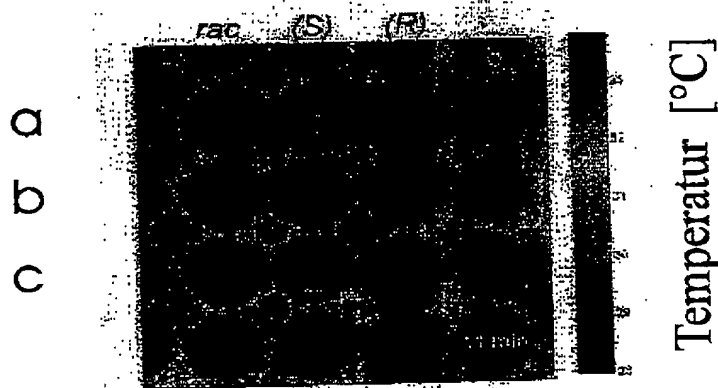


Figure 7/7: IR-image with temperature scale (right) during the enantioselective hydrolysis of various substrates with a cobalt-Salen catalyst at 27 °C

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name. I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought

on the invention entitled Method for Combinatorial Material Development

Using Differential Thermal Images

the specification of which is attached hereto,

or was filed on

as Application Serial No.



I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims.

I acknowledge the duty to disclose information which is material to the patentability of this application in accordance with Title 37, Code of Federal Regulations, §1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, §119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

Prior Foreign Application(s), the priority(ies) of which is/are to be claimed:

197 57 754.7	Germany	12/23/1997
198 26 303.1	Germany	06/12/1998
(Number)	(Country)	(Month/Day/Year Filed)

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose the material information as defined in Title 37, Code of Federal Regulations, §1.56 which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

PCT/EP98/O8214	Dec. 15, 1998	pending
(Application Serial No.)	(Filing Date)	(Status)
		(patented, pending, abandoned)

(Application Serial No.)	(Filing Date)	(Status)
		(patented, pending, abandoned)

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

POWER OF ATTORNEY: As a named Inventor, I hereby appoint the following attorneys and/or agents to prosecute this application and transact all business in the Patent and Trademark Office connected therewith:

Kurt G. Briscoe, Reg. No. 33,141; William C. Gerstenzang, Reg. 27,552; Carmella A. O'Gorman, Reg. No. 33,769
Stephen G. Ryan, Reg. No. 39,015 and Lorimer P. Brooks, all of 660 White Plains Road, Tarrytown, New York
10591, William R. Robinson, Reg. No. 27,224, Davy E. Zoneraich, Reg. No. 37,267 and Mark A. Montana, Reg.
44,948, all of 721 Route 202-206, Bridgewater, New Jersey 08807 my attorneys with full power of substitution and
revocation

Send Correspondence To:

NORRIS, McLAUGHLIN & MARCUS
660 White Plains Road, 4th Fl.
Tarrytown, New York 10591-51444

Direct Telephone Calls To:

(914) 332-1700

Full Name Of Sole or First Inventor <u>Wilhelm E. Maier</u>	Inventor's Signature <u>[Signature]</u>	Date 06/16/2000
Residence 45289 Essen <u>DEX</u>	Citizenship German	
Post Office Address Max-Planck-Institut für Kohlenforschung, Kaiser-Wilhelm-Platz 1, 45470 Mülheim an der Ruhr, Germany		
Full Name Of Second Inventor <u>Arnold Holzwarth</u>	Inventor's Signature <u>[Signature]</u>	Date 06/20/2000
Residence 69100 Villeurbanne, France	Citizenship German	
Post Office Address Institut de Recherches sur la Catalyse, GRC, 2 ^e avenue Albert Einstein, F-69626 Villeurbanne, France <u>BR</u>		
Full Name Of Third Inventor	Inventor's Signature	Date
Residence	Citizenship	
Post Office Address		
Full Name Of Fourth Inventor	Inventor's Signature	Date
Residence	Citizenship	
Post Office Address		
Full Name Of Fifth Inventor	Inventor's Signature	Date
Residence	Citizenship	
Post Office Address		
Full Name Of Sixth Inventor	Inventor's Signature	Date
Residence	Citizenship	
Post Office Address		
Full Name Of Seventh Inventor	Inventor's Signature	Date
Residence	Citizenship	
Post Office Address		